# Center for Regulatory Services, Inc.

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**April 22, 2015** 

CBIC Control Number

364818

U.S. Environmental Protection Agency - East Attn: TSCA Section 8(e) Room 6428

1201 Constitution Avenue, NW Washington, DC 20004

SUBJECT:

TSCA 8(e) Notification

LVE

L-03-153

The enclosed aquatic tox results that came the attention of JSR Micro, Inc., April 17, 2015, for the subject substance that is identified in Low Volume Exemption L-03-153.

The results of the aquatic tox testing of the substance is only identified as **BOSNET**.

48-hour EC50 Acute Immobilization in *Daphnia magna* – <1.0 mg/L 72-hour EC50 Algal Growth Inhibition in Pseudokirchneriella subcapitata - <1.0 mg/L

Please feel free to contact the undersigned if you have any questions or if we can provide additional information.

Sincerely,

William A. Olson, Ph.D. Agent JSR Micro, Inc.

WAO:gbt JSR-8E-BOSNET

**Enclosures** 

2 Aquatic Tox Reports (4 pages)

ec: Y. Ueda/ T. Ozag, JSR (w/o Enclosures)



Receipt number	662-14-E-6879
Study number	96879

March 17, 2015

## TEST REPORT

- A 48-hour Acute Immobilization Study in Daphnia magna -

Chemicals Evaluation and Research Institute,

Japan, Kurume

3-2-7, Miyanojin, Kurume-shi,

Fukuoka 839-0801, Japan

1. Test item

BOSNET

2. Sponsor

JSR Corporation

3. Objective

To determine acute effects of the test item to daphnids

4. Dates Exposure initiation

February 23, 2015

Exposure termination

rmination February 25, 2015

5. Materials and methods

Test organism

Daphnia magna (Clone A)

Exposure conditions

Exposure duration:

48 hours

Test type:

Static regime

Test concentration: Preparation of test solution: 100, 10, 1.0 mg/L as nominal concentration, and a control The test item and dilution water were mixed to prepare each nominal concentration and stirred for 48 hours under shading. Then the suspension was filtered with a glass fiber filter (GB-140, 0.4 µm pore size, Toyo Roshi) by suction to prepare the test

solution. The test item was treated under yellow fluorescent light.

**Environmental conditions** 

Dilution water:

Dechlorinated tap water

Temperature:

20±1°C

Number of organisms:

20 daphnids/test level (5 daphnids/test vessel, 4 replicates)

Volume of test solution:

400 mL/test level (100 mL/test vessel, 4 replicates)

Test vessel:

100 mL glass beaker

Lighting condition:

Shading condition

It was conducted under the yellow fluorescent light at the preparation of test item, handing of test organism, measurement of water quality and observation of test organisms, and under the

room light at filtering the test solutions.

Feeding:

No feeding

Aeration:

No aeration

Observation and measurements

Observation of organisms:

Immobility was observed at 24 and 48 hours after exposure.

Daphnids were considered immobile if they were not able to swim

within 15 seconds after gentle agitation of the test vessel.

Water quality:

Dissolved oxygen concentration and pH were measured of 100

mg/L and the control at the start and end of exposure.

Appearance of test solution:

Colorless and clear (at the start of exposure: visual)

### 6. Result

48-hour median effective concentration (48hr EC<sub>50</sub>): <1.0 mg/L (nominal concentration)

Table Result of immobility and quality of test solution

Test level (mg/L)	Immobility (%)		Dissolved oxygen concentration (mg/L)		рН	
	24 hours	48 hours	At the start	At the end	At the start	At the end
Control	0	0	8.9	8.9	7.8	7.8
1.0	90	100				
10	100	100				
100	100	100	8.8	8.9	7.8	7.7



Receipt number	662-14-E-6878
Study number	96878

March 17, 2015

# TEST REPORT

Algal Growth Inhibition Study in Pseudokirchneriella subcapitata

Chemicals Evaluation and Research Institute, Japan, Kurume 3-2-7, Miyanojin, Kurume-shi, Fukuoka 839-0801, Japan

1. Test item

**BOSNET** 

2. Sponsor

JSR Corporation

3. Objective

To determine the effects of the test item on growth of algae

4. Dates Exposure initiation Exposure termination February 20, 2015 February 23, 2015

5. Materials and methods Test organism

Pseudokirchneriella subcapitata

Exposure conditions

Exposure duration:

Test concentration:

72 hours

Type test:

Incubation with shaking (approximately 100 rpm) 100, 10, 1.0 mg/L as nominal concentration and a control

Preparation of test solution:

The test item and medium were mixed to prepare each nominal concentration and stirred for 48 hours under shading. Then the suspension was filtered with a glass fiber filter (GB-140, 0.4 µm pore size, Toyo Roshi) by suction to prepare the test solution.

The test item was treated under yellow fluorescent light.

**Environmental conditions** 

Medium:

OECD medium

Temperature:

 $21-24^{\circ}$ C (not varied more than  $\pm 2^{\circ}$ C)

Initial cell concentration:

10<sup>4</sup> cells/mL

Volume of test solution:

300 mL/test level (100 mL/test vessel × 3 replicates)

Test vessel:

Sterilized 300 mL Erlenmeyer flask with gas-permeable

silicon rubber plug

Lighting condition:

Nominal 90 μmol·m<sup>-2</sup>·s<sup>-1</sup>

(within  $\pm$  20% of nominal, within  $\pm$  15% from the average

light intensity)

Continuous illumination provided by fluorescent lights

with wavelength range of 400-700 nm

Measurements

Biomass:

Chlorophyll fluorescence value was measured.

Condition of test solution:

pH of 100 mg/L and control were measured at the start and

end of exposure.

Appearance of test solution: Clear and colorless (at the start of exposure: visual)

#### 6. Result

72-hour median effective concentration (72hr  $E_rC_{50}$ ) [Based on growth rate (0-3d)] : <1.0 mg/L (nominal concentration)

No Observed Effect Concentration (NOEC): <1.0 mg/L (nominal concentration)

Table Growth inhibition rate and pH of test solution

Test level	Growth inhibition rate (%)	pН		
(mg/L)	(Growth rate 0-3d)	At the start	At the end	
Control	-	7.9	7.8	
1.0	106	-		
10	136	-	_	
100	159	7.9	7.8	

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